PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

Perry Johnson Laboratory Accreditation, Inc. has assessed the Organization of:

Eurofins BioPharma Product Testing ENCO, Inc.

4810 Executive Park Court, Suite 110, Jacksonville, FL 32216

and hereby declares that the Organization is accredited in accordance with the recognized International Standard:

ISO/IEC 17025:2017

Whereby, technical competence has been confirmed for the associated scope supplement, in the fields of:

Chemical and Dimensional Inspection Testing

(As detailed in the supplement)

Accreditation claims for conformity assessment activities shall only be made from the addresses referenced within this certificate and shall apply solely to those activities identified in the related scope. This Accreditation is granted subject to the Accreditation Body rules governing the Accreditation referred to above, and the Organization hereby commits to observing and complying with those rules in their entirety.

For PJLA:

Tracy Szerszen
President

Perry Johnson Laboratory
Accreditation, Inc. (PJLA)
755 W. Big Beaver, Suite 1325
Troy, Michigan 48084

Initial Accreditation Date:	*Issue Date:*	*Expiration Date:*
January 27, 2026	January 27, 2026	January 27, 2028

Accreditation No.:	*Certificate No.:*
79037	L26-51

The validity of this certificate is maintained through ongoing assessments based on a continuous accreditation cycle. The validity of this certificate should be confirmed through the PJLA website: www.pjlabs.com

Certificate of Accreditation: Supplement

Eurofins BioPharma Product Testing ENCO, Inc.

4810 Executive Park Court, Suite 110, Jacksonville, FL 32216
Contact Name: Cheryl Loze Phone: 904-296-2335

Accreditation is granted to the facility to perform the following conformity assessment activities:

FIELD OF TEST	ITEMS, MATERIALS, OR PRODUCTS TESTED	COMPONENT, CHARACTERISTIC, PARAMETER TESTED	SPECIFICATION OR STANDARD METHOD	TECHNOLOGY OR TECHNIQUE USED	FLEX CODE	LOCATION OF ACTIVITY
Chemical	Raw Materials, Finished Product, Medical Devices	Chemical Safety Assessment/Chemical; Characterization, Extractables and Leachables	ISO 10993-18	GC/MS ICP/MS	F1, F2, F3	F
Chemical	Raw Materials, Finished Product, Medical Devices	Chromatography	USP <621>	HPLC, TLC, GC, IC	F1, F2, F3	F
Chemical	Raw Materials, Finished Product, Medical Devices	Elemental Impurities	USP <232> USP <233>	ICP/MS	F1, F2, F3	F
Chemical	Raw Materials, Finished Product, Medical Devices	Ethylene Oxide Residuals	ISO 10993-7	GC FID	F1, F2	F
Chemical	Raw Materials, Finished Product, Medical Devices	ID of Polymers	USP <197>	FTIR and ATR attachment	F1, F2, F3	F
Chemical	Raw Materials, Finished Product, Medical Devices	Plastic Packaging System	USP <661.1>	FTIR Titration	F1, F2, F3	F
Chemical	Raw Materials, Finished Product, Medical Devices	Plastic Packaging System	USP <661.2>	Visual Appearance	F1, F2, F3	F
Chemical	Raw Materials, Finished Product, Medical Devices	Total Organic Carbon,	USP <643>	TOC Analyzer	F1, F2	F
Chemical	Raw Materials, Finished Product, Medical Devices	UV/Vis Spectroscopy	USP <857>	UV/Vis Spectroscopy	F1, F2, F3	F

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Chemical	Raw Materials, Finished Product, Medical Devices	Thermal Analysis	USP <891>	Differential Scanning Calorimeter	F1, F2	F
Chemical	Raw Materials, Finished Product, Medical Devices	Water Content	USP <921>	Karl Fischer Titration	F1, F2	F
Dimensional Inspection	Raw Materials, Finished Product, Medical Devices	Measurements	ENCO-ATM-032	Dimensional Analysis	F1, F4	F

1. Location of activity:

Location	Location
F	Conformity assessment activity is performed at the CABs fixed facility

2. Flex Code:
   F0- Fixed scope item. No deviations allowed to the line item as identified, except for updating to the most recent version of an accredited standard method after verification.
   F1- Laboratory has the capability to test a new item, material, matrix, or product similar in composition to item, material, matrix, or product identified on the scope
   F2- Laboratory has the capability to introduce the newest revision of an accredited authoritative standard method (with no modifications) identified on the scope
   F3- Laboratory has the capability to introduce a parameter/component/analyte to an accredited test method identified on the scope
   F4- Laboratory has the capability to introduce a new revision of an accredited non-standard method using the same technology or technique identified on the scope
   F5- Laboratory has the capability to introduce a validated method that is equivalent to an accredited method (using same technology or technique) identified on the scope